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NEWS	1			Web Page for STN Seminar Schedule - N. America
NEWS	2	OCT	02	CA/CAplus enhanced with pre-1907 records from Chemisches Zentralblatt
NEWS	3	OCT	19	BEILSTEIN updated with new compounds
NEWS		NOV		Derwent Indian patent publication number format enhanced
NEWS		NOV		WPIX enhanced with XML display format
NEWS		NOV		ICSD reloaded with enhancements
NEWS		DEC		LINPADOCDB now available on STN
NEWS		DEC		BEILSTEIN pricing structure to change
NEWS		DEC		USPATOLD added to additional database clusters
NEWS				IMSDRUGCONF removed from database clusters and STN
NEWS		DEC		DGENE now includes more than 10 million sequences
NEWS		DEC		TOXCENTER enhanced with 2008 MeSH vocabulary in
MEMO	12	DEC	1 /	MEDLINE segment
NEWS	13	DEC	17	MEDLINE and LMEDLINE updated with 2008 MeSH vocabulary
NEWS		DEC		CA/CAplus enhanced with new custom IPC display formats
NEWS		DEC		STN Viewer enhanced with full-text patent content
NEWS	13	DEC	1/	from USPATOLD
MENTO	10	7737	00	
NEWS		JAN		STN pricing information for 2008 now available
NEWS	1/	JAN	10	CAS patent coverage enhanced to include exemplified prophetic substances
NEWS	1.0	JAN	20	USPATFULL, USPAT2, and USPATOLD enhanced with new
NEWS	10	UMIN	20	custom IPC display formats
NEWS	10	JAN	20	MARPAT searching enhanced
NEWS		JAN		USGENE now provides USPTO sequence data within 3 days
NEWS	20	UAIN	20	of publication
NEWS	21	JAN	20	TOXCENTER enhanced with reloaded MEDLINE segment
NEWS		JAN		MEDLINE and LMEDLINE reloaded with enhancements
NEWS				STN Express, Version 8.3, now available
NEWS				PCI now available as a replacement to DPCI
NEWS				IFIREF reloaded with enhancements
NEWS				IMSPRODUCT reloaded with enhancements
NEWS	27	FEB	29	WPINDEX/WPIDS/WPIX enhanced with ECLA and current
				U.S. National Patent Classification
NEWS	28	MAR	31	IFICDB, IFIPAT, and IFIUDB enhanced with new custom
N1111110	0.0		2.0	IPC display formats
NEWS	29	MAR	31	CAS REGISTRY enhanced with additional experimental
NEWS	2.0	1/2 D	2.1	spectra
NEWS	30	MAR	31	CA/CAplus and CASREACT patent number format for U.S.
			0.0	applications updated
NEWS		MAR		LPCI now available as a replacement to LDPCI
NEWS	32	MAR	31	EMBASE, EMBAL, and LEMBASE reloaded with enhancements
NELLO	EVE	2000	DDD	NAME OF STREET, STANSON STREET, TO US 2
NEWS	EXP	KESS		RUARY 08 CURRENT WINDOWS VERSION IS V8.3,

AND CURRENT DISCOVER FILE IS DATED 20 FEBRUARY 2008

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0.21

TOTAL ENTRY SESSION

0.21

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FULL ESTIMATED COST

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STRUCTURE FILE UPDATES: 1 APR 2008 HIGHEST RN 1011527-65-3 DICTIONARY FILE UPDATES: 1 APR 2008 HIGHEST RN 1011527-65-3

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Uploading C:\Program Files\Stnexp\Queries\10 series\10553151\10553151a.str

ring nodes:
1 2 3 4 5 6 7 8 9 10 11 12 13 17 26 27 28 29 30
chain bonds:
1-14 4-15 11-16 12-31 13-18 16-17 18-19 19-20 20-21 31-32
ring bonds:
1-2 1-6 2-3 2-7 3-4 3-10 4-5 5-6 5-11 6-13 7-8 8-9 9-10 11-12 12-13
17-26 17-30 26-27 27-28 28-29 29-30
exact/norm bonds:
1-2 1-6 1-14 3-4 4-5 4-15 5-6 5-11 6-13 11-12 11-16 12-13 13-18 19-20
20-21
exact bonds:
12-31 16-17 18-19 31-32
normalized bonds:
2-3 2-7 3-10 7-8 8-9 9-10 17-26 17-30 26-27 27-28 28-29 29-30

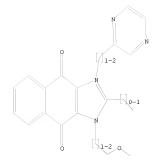
Match level:
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:Atom
1:Atom 12:Atom 13:Atom 14:CLASS 15:CLASS 16:CLASS 17:Atom 18:CLASS
19:CLASS 20:CLASS 21:CLASS 26:Atom 27:Atom 28:Atom 29:Atom 30:Atom 31:CLASS 32:CLASS

L1 STRUCTURE UPLOADED

=> d 11 L1 HAS NO ANSWERS L1 ST

chain nodes :

14 15 16 18 19 20 21 31 32



Structure attributes must be viewed using STN Express query preparation.

=> s 11

SAMPLE SEARCH INITIATED 11:17:01 FILE 'REGISTRY' SAMPLE SCREEN SEARCH COMPLETED -0 TO ITERATE

100.0% PROCESSED 0 ITERATIONS 0 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

COMPLETE BATCH PROJECTED ITERATIONS: 0 TO

PROJECTED ANSWERS: 0 TO

0 SEA SSS SAM L1 L2

=> s l1 full

FULL SEARCH INITIATED 11:17:05 FILE 'REGISTRY'

FULL SCREEN SEARCH COMPLETED -5 TO ITERATE

100.0% PROCESSED 5 ITERATIONS

5 ANSWERS

SEARCH TIME: 00.00.01

L3 5 SEA SSS FUL L1

=> d scan

L3 5 ANSWERS REGISTRY COPYRIGHT 2008 ACS on STN

1H-Naphth[2,3-d]imidazolium, 4,9-dihydro-1-(2-methoxyethyl)-2-methyl-4,9dioxo-3-(2-pyrazinylmethyl)-, bromide (1:1)

MF C20 H19 N4 O3 . Br

• Br-

ONE OR MORE TAUTOMERIC DOUBLE BONDS NOT DISPLAYED IN THE STRUCTURE

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):4

- L3 5 ANSWERS REGISTRY COPYRIGHT 2008 ACS on STN
- IN 1H-Naphth[2,3-d]imidazolium, 4,9-dihydro-1-(2-methoxyethyl)-2-methyl-4,9-dioxo-3-(pyrazinylmethyl)- (9CI)
- MF C20 H19 N4 O3
- CI COM

ONE OR MORE TAUTOMERIC DOUBLE BONDS NOT DISPLAYED IN THE STRUCTURE

- L3 5 ANSMERS REGISTRY COPYRIGHT 2008 ACS on STN
 IN H=Naphth[2,3-d]imidazollum, 4,9-dihydro-1-(2-methoxyethyl)-2-methyl-4,9dioxo-3-(pvrazinvlmethyl)-, chloride (9CI)
- MF C20 H19 N4 O3 . C1

● C1-

ONE OR MORE TAUTOMERIC DOUBLE BONDS NOT DISPLAYED IN THE STRUCTURE

- L3 5 ANSWERS REGISTRY COPYRIGHT 2008 ACS on STN
- IN 1H-Naphth[2,3-d]imidazolium, 4,9-dihydro-1-(2-methoxyethy1)-2-methy1-3-[(5-
- methylpyrazinyl)methyl]-4,9-dioxo- (9CI) MF C21 H21 N4 O3
- CI COM

Me

ONE OR MORE TAUTOMERIC DOUBLE BONDS NOT DISPLAYED IN THE STRUCTURE

- MF C21 H21 N4 O3 . C1

● c1=

ONE OR MORE TAUTOMERIC DOUBLE BONDS NOT DISPLAYED IN THE STRUCTURE

ALL ANSWERS HAVE BEEN SCANNED

=> file caplus COST IN U.S. DOLLARS

ARS SINCE FILE
ENTRY
ST 180.20

TOTAL

SESSION

180.41

FULL ESTIMATED COST

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=> s 13 L4 6 L3

=> d 14 1-6 ibib hitstr

L4 ANSWER 1 OF 6 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2008:252630 CAPLUS DOCUMENT NUMBER: 148:269467

TITLE: Stabilized pharmaceutical composition of

naphthoimidazolium derivative

Murai, Makoto; Yonemochi, Yuichi; Shakushiro, Kohsuke; INVENTOR(S):

Kasai, Akihiro

PATENT ASSIGNEE(S): Astellas Pharma Inc., Japan SOURCE: PCT Int. Appl., 24pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

	PATENT :	KIND DATE				APPL	ICAT	DATE											
	WO 2008023807					A1		20080228		WO 2007-JP66495					20070824				
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		BY,	KG,	KZ,	MD,	RU,	TJ,	TM											
PRIO	RITY APP	LN. :	INFO	. :						JP 2	006-	2292	03	- 1	A 2	0060	825		

RN

A 20070823

WO 2007-JP66340

781661-94-7 IT RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(stabilized pharmaceutical composition of naphthoimidazolium derivative) 781661-94-7 CAPLUS

CN 1H-Naphth[2,3-d]imidazolium, 4,9-dihydro-1-(2-methoxyethy1)-2-methy1-4,9dioxo-3-(2-pyrazinylmethyl)-, bromide (1:1) (CA INDEX NAME)

● Br-

ONE OR MORE TAUTOMERIC DOUBLE BONDS NOT DISPLAYED IN THE STRUCTURE REFERENCE COUNT: 3.0 THERE ARE 30 CITED REFERENCES AVAILABLE FOR THIS RECORD, ALL CITATIONS AVAILABLE IN THE RE FORMAT L4 ANSWER 2 OF 6 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2007:1451296 CAPLUS

DOCUMENT NUMBER: 148:228885

TITLE: YM-155: apoptosis inducer survivin expression

inhibitor oncolvtic AUTHOR(S):

Wang, Y.; Serradell, N.; Bolos, J.; Rosa, E. CORPORATE SOURCE: IBC, Frederick, MD, 21702, USA

SOURCE: Drugs of the Future (2007), 32(10), 879-882

CODEN: DRFUD4; ISSN: 0377-8282 PUBLISHER: Prous Science

DOCUMENT TYPE:

Journal: General Review

LANGUAGE: English ΤТ

781661-94-7 RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL

(Biological study); USES (Uses) (YM-155 monotherapy showed promising anticancer activity and acceptable toxicity profile in patient with cancer)

RM 781661-94-7 CAPLUS

1H-Naphth[2,3-d]imidazolium, 4,9-dihydro-1-(2-methoxyethy1)-2-methy1-4,9-CN dioxo-3-(2-pyrazinylmethyl)-, bromide (1:1) (CA INDEX NAME)

Br -

ONE OR MORE TAUTOMERIC DOUBLE BONDS NOT DISPLAYED IN THE STRUCTURE

ANSWER 3 OF 6 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2007:996056 CAPLUS

DOCUMENT NUMBER: 147:461711

TITLE: YM155, a Novel Small-Molecule Survivin Suppressant,

Induces Regression of Established Human

Hormone-Refractory Prostate Tumor Xenografts

Nakahara, Takahito; Takeuchi, Masahiro; Kinoyama, Isao; Minematsu, Tsuyoshi; Shirasuna, Kenna;

Matsuhisa, Akira; Kita, Aya; Tominaga, Fumiko;

Yamanaka, Kentaro; Kudoh, Masafumi; Sasamata, Masao Institute for Drug Discovery Research, Astellas CORPORATE SOURCE:

Pharma, Inc., Tsukuba-shi, Ibaraki, Japan

SOURCE: Cancer Research (2007), 67(17), 8014-8021 CODEN: CNREA8; ISSN: 0008-5472

PUBLISHER: American Association for Cancer Research

DOCUMENT TYPE: Journal

LANGUAGE: English

781661-94-7, YM 155

AUTHOR(S):

RL: DMA (Drug mechanism of action); PAC (Pharmacological activity); PKT

(Pharmacokinetics); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(YM155, a novel small-mol. survivin suppressant, induces regression of established human hormone-refractory prostate tumor xenografts)

RN 781661-94-7 CAPLUS

CN

1H-Naphth[2,3-d]imidazolium, 4,9-dihydro-1-(2-methoxyethyl)-2-methyl-4,9-dioxo-3-(2-pyrazinylmethyl)-, bromide (1:1) (CA INDEX NAME)

• Br-

ONE OR MORE TAUTOMERIC DOUBLE BONDS NOT DISPLAYED IN THE STRUCTURE
REFERENCE COUNT: 37 THERE ARE 37 CITED REFERENCES AVAILABLE FOR THIS

RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 4 OF 6 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2004:902370 CAPLUS

DOCUMENT NUMBER: 141:379945

TITLE: Process for preparation of imidazolium bromide

derivative and its crystals
INVENTOR(S): Kinoyama, Isao; Sakamoto, Kenichirou; Okui, Hiroki;

Hamada, Noritaka; Matsuhisa, Akira

PATENT ASSIGNEE(S): Yamanouchi Pharmaceutical Co., Ltd., Japan

SOURCE: PCT Int. Appl., 23 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.					KIND DATE				APPLICATION NO.							DATE			
WO 2004092160					A1	_	20041028			WO 2	004-		20040414						
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		TD,	TG																
CA 2522486							2004	1028		CA 2	004-	20040414							

JP	2004	A	A 20041125				JP 2004-118511					20040414							
JP	3618341				B2 20050209														
EP	1614686			A1	20060111			E	7274	20040414									
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CN	1791	595			A		2006	0621	C	N 20	04-1	3001	3226		2	0040	414		
MX	2005	PA11	078		A	- 2	2006	0124	M	X 20	05-I	PA11	078		2	0051	014		
US	2006	0223	831		A1	- 1	2006	1005	U	S 20	05-5	5531.	51		2	0051	014		
IN	2005	DN04	719		A	- 1	2007	0105	I	N 20	05-1	DN47	19		2	0051	017		
PRIORITY	APP	LN.	INFO	. :					J.	P 20	03-	1097	93		A 2	0030	415		
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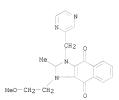
OTHER SOURCE(S): ΙT

CASREACT 141:379945 781661-94-7P

RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation) (preparation of imidazolium bromide derivative and its crystals)

RN 781661-94-7 CAPLUS

CN 1H-Naphth[2,3-d]imidazolium, 4,9-dihydro-1-(2-methoxyethy1)-2-methy1-4,9dioxo-3-(2-pyrazinylmethyl)-, bromide (1:1) (CA INDEX NAME)



Br -

ONE OR MORE TAUTOMERIC DOUBLE BONDS NOT DISPLAYED IN THE STRUCTURE

355406-09-6P

RL: SPN (Synthetic preparation); PREP (Preparation)

(preparation of imidazolium bromide derivative and its crystals)

RN 355406-09-6 CAPLUS

CN 1H-Naphth[2,3-d]imidazolium, 4,9-dihydro-1-(2-methoxyethy1)-2-methy1-4,9dioxo-3-(pyrazinylmethyl)-, chloride (9CI) (CA INDEX NAME)

c1 =

ONE OR MORE TAUTOMERIC DOUBLE BONDS NOT DISPLAYED IN THE STRUCTURE

REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS
RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L4 ANSWER 5 OF 6 CAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2003:349225 CAPLUS

DOCUMENT NUMBER: 2003:34922.

TITLE: Pharmaceutical compositions containing condensed

imidazolium derivatives

INVENTOR(S): Matsuhisa, Akira; Kinoyama, Isao; Toyoshima, Hiroshi;

Nakahara, Takato; Takeuchi, Masahiro; Okada, Minoru

PATENT ASSIGNEE(S): Yamanouchi Pharmaceutical Co., Ltd., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 27 pp.

CÔDEN: JKXXAF
DOCUMENT TYPE: Patent

LANGUAGE: Japanese FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2003128548	A	20030508	JP 2002-230691	20020808
JP 3634328	B2	20050330		
PRIORITY APPLN. INFO.:			JP 2001-243397 A	20010810

OTHER SOURCE(S): MARPAT 138:348689

T 355406-09-6P 355406-23-4P

RL: ADV (Adverse effect, including toxicity); PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of condensed imidazolium derivs. for anticancer agents) RN 355406-09-6 CAPLUS

CN 1H-Naphth[2,3-d]imidazolium, 4,9-dihydro-1-(2-methoxyethyl)-2-methyl-4,9-dioxo-3-(pyrazinylmethyl)-, chloride (9CI) (CA INDEX NAME)

c1=

ONE OR MORE TAUTOMERIC DOUBLE BONDS NOT DISPLAYED IN THE STRUCTURE RN 355406-23-4 CAPLUS

1H-Naphth[2,3-d]imidazolium, 4,9-dihydro-1-(2-methoxyethyl)-2-methyl-3-[(5-CN methylpyrazinyl)methyl]-4,9-dioxo-, chloride (9CI) (CA INDEX NAME)

C1 =

ONE OR MORE TAUTOMERIC DOUBLE BONDS NOT DISPLAYED IN THE STRUCTURE

L4 ANSWER 6 OF 6 CAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2001:617983 CAPLUS

DOCUMENT NUMBER: 135:180768

TITLE:

Preparation of naphth[2,3-d]imidazolium halides, naphth[2,3-d]imidazoles and aminonaphthalene-1,4-dione derivatives as antitumor agents

INVENTOR(S): Matsuhisa, Akira; Kinoyama, Isao; Toyoshima, Akira; Nakahara, Takahito; Takeuchi, Masahiro; Okada, Minoru

PATENT ASSIGNEE(S): Yamanouchi Pharmaceutical Co., Ltd., Japan

SOURCE: PCT Int. Appl., 61 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: Japanese

PA	PATENT NO.						DATE		APPLICATION NO.						DATE			
WO	2001060803			A1 20010823			WO 2001-JP1036						20010214					
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		SE,	SG,	SI,	SK,	SL,	TJ,	TM,	TR,	T	r, TZ,	UA,	UG,	US,	UZ,	VN,	YU,	
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	2395		A1										20010214					
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EP	1256	576							EP 2001-904433						2	0010	214	
EP	1256	576			B1 20050615													
	R:	ΑT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GE	R, IT,	LI,	LU,	NL,	SE,	MC,	PT,	
		IE,	SI,	LT,	LV,	FI,	RO,	MK,	CY,	ΑI	, TR							
JP	3397	320			B2		2003	0414		JP	2001-	5601	88		2	0010	214	
AT	2979	01			T		2005	0715		ΑT	2001-	9044	33		2	0010	214	
PT	1256	576			T		2005	0930		PΤ	2001-	9044	33		2	0010	214	
	2243				Т3						2001-							
US	2003	0114	508		A1		2003	0619		US	2002-	1694	09		2	0020	701	
US	6734	203			B2		2004	0511										
PRIORIT	Y APP	LN.	INFO	. :						JΡ	2000-	3664	5		A 2	0000	215	
										JΡ	2000-	2614	89		A 2	0000	830	
										WO	2001-	JP10	36		W 2	0010	214	
OTHER S	HER SOURCE(S).					TAG	135.	1807	5.8									

OTHER SOURCE(S): MARPAT 135:180768

IT 355406-09-6P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of naphth[2,3-d]imidazolium halides and aminonaphthalene-1,4dione derivs. as antitumor agents)

RN 355406-09-6 CAPLUS

CN 1H-Naphth[2,3-d]imidazolium, 4,9-dihydro-1-(2-methoxyethyl)-2-methyl-4,9-dioxo-3-(pyrazinylmethyl)-, chloride (9CI) (CA INDEX NAME)

IT 355406-23-4P

RL: SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of naphth[2,3-d]imidazolium halides and aminonaphthalene-1,4-dione derivs. as antitumor agents)

RN 355406-23-4 CAPLUS

CN 1H-Naphth[2,3-d]imidazolium, 4,9-dihydro-1-(2-methoxyethyl)-2-methyl-3-[(5-methylpyrazinyl)methyl]-4,9-dioxo-, chloride (9CI) (CA INDEX NAME)

● C1-

ONE OR MORE TAUTOMERIC DOUBLE BONDS NOT DISPLAYED IN THE STRUCTURE
REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS
RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

=>

=> log hold COST IN U.S. DOLLARS

FULL ESTIMATED COST

SINCE FILE TOTAL ENTRY SESSION 34.98 215.39

SESSION WILL BE HELD FOR 120 MINUTES

STN INTERNATIONAL SESSION SUSPENDED AT 11:35:20 ON 02 APR 2008